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REMARKS

With entry of the present amendment, the application will contain claims 1-10, all of which are under examination.

Allowable Subject Matter

The Examiner is thanked for the allowance of claim 5 as specified in the last Office Action, page 6, paragraph 16.

Priority

On page 2, paragraph 2, of the Office Action, the Examiner indicates that the certified translation is not currently part of the case record. This certified translation was first timely filed on July 2, 2003, as shown by the mailroom date stamp on the postcard receipt attached hereto as Exhibit A. A second copy of the certified translation was next timely filed on July 11, 2003. Notwithstanding two filings, the Examiner later stated: "...this translation document has not been received; hence it is not currently part of the record." (Office Action mailed September 16, 2003; page 2, line 7). It is undisputed that the Examiner did not receive it. It is also undisputed that the PTO did receive it. Since the PTO did receive it, it is part of the record.

It is incumbent upon the PTO to locate the document and make it a part of the record.

Notwithstanding the fact that the certified translation has already been twice submitted, it is submitted concurrently herewith a third time for the convenience of the Examiner and in order to advance prosecution. This third submission is labeled Exhibit B.

The AZECHI Declaration

The Examiner has not commented on a certain four-page document entitled "DECLARATION" signed by the inventor, AZECHI, on May 30, 2003 (the AZECHI Declaration). The AZECHI Declaration was filed concurrently with the prior filed certified translations as shown by Exhibit A. An additional copy of the AZECHI Declaration is filed concurrently herewith as Exhibit C in case the earlier filed copies have not found their way to the desk of the Examiner.

Issues Under 35 USC § 112, First Paragraph

The rejection of claims 1-3 and 6-9 in paragraph 4 of the last Office Action is traversed but has been rendered moot by the present amendments. As presently amended, all claims are

limited to thermoplastic resins. The mold, where recited, is indicated as being a metal mold.

The rejection in paragraph 5 of claims 1-3 and 6-9 is traversed but has been rendered moot by the present amendments.

The rejection of claim 3 in paragraph 6 of the last Office Action is traversed but has been rendered moot by the present amendment. As presently amended, the property there recited is indicated as having been measured with steel as the metal.

The rejection in paragraph 7 of claims 1-3 and 6-9 is traversed but has been overcome by the present amendments. As presently amended, the claims are limited to "thermoplastic resins" and to metal molds.

Issues Under 35 USC § 112, Second Paragraph

The rejection of claims 1-3 and 6-9 is traversed but has been overcome by the present amendments. The Examiner is thanked for the suggested amendment to claim 1. That amendment has been made. With this amendment, the objection to claim 1 is believed to have been overcome. The other

claims there rejected are dependent on claim 1 and are, therefore, now free of objection.

It is believed that the amendments to claim 6 now provide clear antecedent basis.

Issues Under 35 USC § 102/103

These grounds of rejection have been overcome by the prior and the concurrent filing of the certified translation for the reasons mentioned beginning in the last paragraph on page 7 of the document entitled "Reply Under 37 CFR § 1.111" filed on or about July 3, 2003.

CONCLUSION

Should there be any outstanding matters that need to be resolved in the present application, the Examiner is respectfully requested to contact David R. Murphy (Reg. No. 22,751) at the telephone number of the undersigned below, to conduct an interview in an effort to expedite prosecution in connection with the present application.

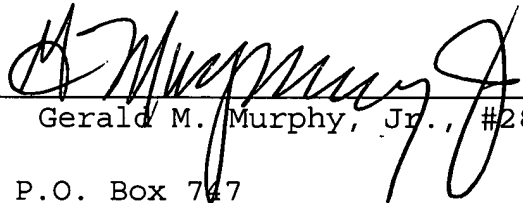
If necessary, the Commissioner is hereby authorized in this, concurrent, and future replies, to charge payment or credit any overpayment to Deposit Account No. 02-2448 for any

Appl. No.: 10/035,213

additional fees required under 37 C.F.R. §§ 1.16 or 1.17;
particularly, extension of time fees.

Respectfully submitted,

BIRCH, STEWART, KOLASCH & BIRCH, LLP

By 
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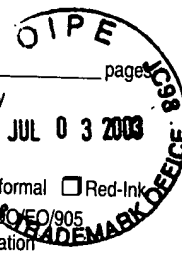
0171-0808P
GMM/DRM/enm/drm


Exhibits:

- A. Postcard receipt bearing mail room date stamp of July 3, 2003, proving filing of certified translation.
- B. Third copy of certified translation consisting of:
 - (1) Certification (1 page)
 - (2) Translation of cover sheet (3 pages)
 - (3) Specification (30 pages)
 - (4) Drawings (6 pages)
 - (1) Abstract (1 page)
- C. Third copy of document entitled "DECLARATION" signed by the inventor, AZECHI, on May 30, 2003 (4 pages)

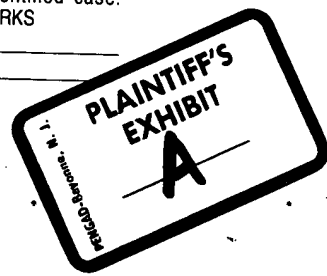
Papers Filed herewith on: July 3, 2003
DOCKET NO.: 0171-0808P ATTY.: GMM/DRM
APPLICANT(S): Syuuichi AZECHI et al.
APPLN. NO: 10/635,213 FILED: January 4, 2002
PAT NO.:

- ☐ New Application with Transmittal Letter
☐ Utility ☐ Design ☐ CIP ☐ PCT ☐ Provisional
☐ Filing Under 37 CFR 1.53(b) ☐ CONT ☐ DIV
☐ Filing Under 37 CFR 1.53(d) (CPA)
☐ Filing Under 37 CFR 1.114(RCE)
☐ Specification Consisting of: _____ pages
☐ Combined Declaration & Power of Attorney
☐ Assignment / Cover Letter
☐ Letter to Official Draftsman
☐ Drawings _____ Sheets ☐ Formal ☐ Informal ☐ Red-Ink
☐ Completion of Filing Requirements, PCT/RO/EO/905
or Formalities Letter and Executed Declaration
☐ Priority Document(s) / Cover Letter, No. Doc. _____
☒ Amendment: 37 CFR 1.111
☒ Transmtl Ltr ☒ Large Entity ☐ Small Entity
☐ Response
☐ Information Discl Stmt. PTO-1449(s) _____ doc(s)
☐ Notice of Appeal ☐ Appeal Brief
☐ Issue Fee Transmittal ☐ Sequence Listing
☐ FEES: _____
☐ Letter: dt 5/30/03
☒ Other: DECLARATION by AZECHI
CERTIFIED TRANSLATION



DOCKET NO. 0171-0808P

Receipt is hereby acknowledged of the papers filed as indicated in connection with the above identified case.
COMMISSIONER OF PATENTS AND TRADEMARKS
Due Date: July 4, 2003
Handcarry: _____



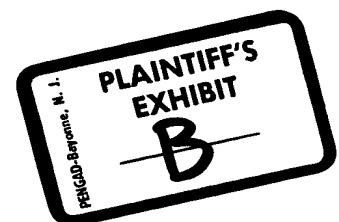
COPY

C E R T I F I C A T I O N

I, Takashi KOJIMA of Ginza Ohtsuka Bldg., 2F, 16-12, Ginza 2-chome, Chuo-ku, Tokyo, Japan, hereby certify that I am the translator of the accompanying certified official copy of the documents in respect of an application for a patent filed in Japan on the 5th of January, 2001 and of the official certificate attached thereto, and certify that the following is a true and correct translation to the best of my knowledge and belief.

Dated this 5th day of June, 2003

Takashi Kojima
Takashi KOJIMA



(Translation)

PATENT OFFICE
JAPANESE GOVERNMENT

This is to certify that the annexed is a true copy of the following application as filed with this Office.

Date of Application: January 5, 2001

Application Number: Japanese Patent Application
No. 2001-000536

Applicant(s): Shin-Etsu Chemical Co., Ltd.

August 24, 2001

Commissioner,
Patent Office

Kozo OIKAWA
(sealed)

Certificate No. 2001-3075978

2001-000536

[Paper]	APPLICATION FOR PATENT
[Reference Number]	12729
[Application Date]	January 5, 2001
[Destination]	The Commissioner of the Patent Office Kozo OIKAWA
[International Patent Classification]	C08L 83/07
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[Patent Attorney]	
[Name]	Yuko NISHIKAWA

[Official Fee]

[Deposit Account Number] 003207

[Amount of Fee] 21000

[List of Documents Attached]

[Document] Specification 1

[Document] Drawing 1

[Document] Abstract 1

[Necessity of Proof] Yes

2001-000536

[SPECIFICATION]

[TITLE OF THE INVENTION] Silicone Rubber Adhesive
Composition and Integrally Molded
Article of Silicone Rubber and
Thermoplastic Resin

[CLAIMS]

[Claim 1] A silicone rubber adhesive composition
comprising

10 (A) 100 parts by weight of a heat curable
organopolysiloxane composition,

 (B) 1 to 100 parts by weight of reinforcing silica
fines, and

 (C) 0.1 to 50 parts by weight of an organic compound
15 or organosilicon compound having an epoxy equivalent of 100
to 5,000 g/mol and containing at least one aromatic ring in
a molecule.

[Claim 2] The silicone rubber adhesive composition of
claim 1 wherein component (C) is an organosilicon compound
20 containing at least one Si-H group in a molecule.

[Claim 3] The silicone rubber adhesive composition of
claim 1 or 2 which provides a greater bond strength to
organic resins than to metals.

[Claim 4] An integrally molded article comprising a
25 silicone rubber adhesive composition in the cured state and
a thermoplastic resin, said silicone rubber adhesive
composition comprising

 (A) 100 parts by weight of a heat curable
organopolysiloxane composition,

30 (B) 1 to 100 parts by weight of reinforcing silica
fines, and

 (C) 0.1 to 50 parts by weight of an organic compound
or organosilicon compound having an epoxy equivalent of 100
to 5,000 g/mol and containing at least one aromatic ring in
35 a molecule.

[Claim 5] The integrally molded article comprising a
silicone rubber adhesive composition in the cured state and

a thermoplastic resin of claim 4 wherein component (C) is an organosilicon compound containing at least one Si-H group in a molecule.

[DETAILED DESCRIPTION OF THE INVENTION]

5 [0001]

[Technical Field of the Invention]

This invention relates to a silicone rubber adhesive composition which can be simply and briefly molded by injection molding, extrusion or mechanical application using
10 a robot, is suited for primerless molding and hence, integral molding with a thermoplastic resin, and exhibits the unique nature that when molded together with a thermoplastic resin in a mold which has been or has not been surface treated as by chromium plating, the composition is
15 adherent only to the resin, but not to the mold. It also relates to an integrally molded article of a cured product (silicone rubber) of a silicone rubber adhesive composition bonded to a thermoplastic resin, which is effective for fields including electric, electronic, automotive, precision
20 machinery and building fields.

[0002]

[Prior Art and Problem to be Solved by the Invention]

Nowadays silicone rubber, due to its excellent characteristics, finds widespread use in a variety of fields
25 including electric, electronic, automotive and building fields. The demands for the silicone rubber in these fields include workability, seal and insulation during assembly. In some applications where the adhesion of silicone rubber to resins is necessary, adhesion is achieved by way of
30 primers. The process requires the step of applying primer and is thus cumbersome. One solution is a self-adhesive type silicone rubber obtained by adding an adhesive to a silicone rubber composition. The self-adhesive type silicone rubber eliminates the application step, shortens
35 the operating time, reduces the cost, and improves the efficiency of operation. It is effective in manufacturing integrally molded articles with resins.

[0003]

In connection with the primerless molding of addition type heat-curable silicone rubber, a number of reports have been made for bonding silicone rubber to organic resins. In one exemplary attempt, self-adhesive type silicone rubber material is cured on resins. A number of proposals have been made relating to specific adhesive components for use in the self-adhesive type silicone rubber. Also, JP-B 2-34311 discloses to add an organopolysiloxane containing at least 30 mol% of hydrogen atoms directly bonded to silicon atoms to an organic resin whereby the resin is bondable to an addition curing type silicone rubber. JP-B 63-4529 discloses the integration of silicone rubber to organic resin by physical engagement. JP-A 63-183843 discloses the integral adhesion of silicone rubber to an olefin resin having grafted thereto a compound having an aliphatic unsaturated group and a silicon atom-bonded hydrolyzable group. JP-A 9-165516 and JP-A 9-165517 propose a process in which a thermoplastic resin having added thereto a compound having unsaturated groups and hydrogen atoms directly bonded to silicon atoms is adhesively joined and integrated with silicone rubber as well as an integrally molded article in which a thermoplastic resin having blended therein a thermoplastic oligomer containing aliphatic unsaturated groups is integrally molded with an oil-bleeding silicone rubber.

[0004]

However, heat curable silicone rubber of the addition type fails to achieve within brief molding a sufficient bond to general thermoplastic resins, fibers and films, such as ABS, PPO, PPS, PC, PE, PP, PBT, acrylic resins, phenolic resins, polyimide resins, PA and aromatic PA and other copolymers in resin, fiber or film form. To acquire a bonding ability, the resins must be modified as suggested in the above patents. Modification of resins requires extra steps and an increased cost, and the modified resins sometimes become susceptible to quality alteration. Even if

a bond is established without resin modification, there still arises a problem that when an integrated silicone rubber/resin article is formed in a mold, the molded article sticks to the mold. In particular, none of available
5 silicone rubber compositions fully satisfy the requirements when combined with polyamide resins and fibers. It remains of interest to develop an integrally molded article of thermoplastic resin and silicone rubber which when formed in a mold, does not stick to the mold and establishes a firm
10 bond between the thermoplastic resin and the silicone rubber.

[0005]

The present invention has responded to the above desire. An object of the invention is to provide a silicone
15 rubber adhesive composition which can be firmly integrated with thermoplastic resins, as well as an integrally molded article of the cured product (silicone rubber) thereof with a thermoplastic resin.

[0006]

20 [Means for Solving the Problem and Embodiment of the Invention]

The present inventors have earnestly studied in order to attain the above object. As a result, it has been found that blending an organopolysiloxane composition of the heat
25 curing type with an organic compound or organosilicon compound containing an amount of epoxy group to give an epoxy equivalent of 100 to 5,000 g/mol and at least one aromatic ring in a molecule affords a silicone rubber adhesive composition which is firmly bondable to
30 thermoplastic resins. When this silicone rubber adhesive composition is molded and joined with a thermoplastic resin, there is obtained an integrally molded article in which the silicone rubber is firmly bonded to the thermoplastic resin. Even under brief curing conditions using an injection
35 molding technique, the silicone rubber adhesive composition develops a sufficient bonding force to the thermoplastic resin. The integrally molded article of silicone

rubber/thermoplastic resin can be released from the mold in a practically acceptable way. The silicone rubber adhesive composition is effective even to highly crystalline polyamide resins and fibers such as nylon 66 and aromatic polyamides.

[0007]

Accordingly, the invention provides a silicone rubber adhesive composition comprising

(A) 100 parts by weight of a heat curable organopolysiloxane composition,

(B) 1 to 100 parts by weight of reinforcing silica fines, and

(C) 0.1 to 50 parts by weight of an organic compound or organosilicon compound having an epoxy equivalent of 100 to 5,000 g/mol and containing at least one aromatic ring in a molecule. In a preferred embodiment, component (C) is an organosilicon compound containing at least one Si-H group in a molecule. The composition most often provides a greater bond strength to organic resins than to metals. Also contemplated herein is an integrally molded article comprising the cured silicone rubber adhesive composition (i.e., silicone rubber) and a thermoplastic resin.

[0008]

The following is the detailed description of the invention.

The silicone rubber adhesive composition of the invention includes as essential components,

(A) a heat curable organopolysiloxane composition,

(B) reinforcing silica fines, and

(C) an organic compound or organosilicon compound having an epoxy equivalent of 100 to 5,000 g/mol and containing at least one aromatic ring in a molecule.

[0009]

Component (A) is a heat curable organopolysiloxane composition which may be either an addition reaction curing type organopolysiloxane composition or an organic peroxide

curing type organopolysiloxane composition, with the former being preferred.

The addition reaction curing type organopolysiloxane composition is preferably defined as comprising

- 5 (1) 100 parts by weight of an organopolysiloxane having on the average at least two alkenyl groups in a molecule,
(2) 0.1 to 50 parts by weight of an organohydrogenpolysiloxane having on the average at least two hydrogen atoms attached to silicon atoms in a molecule,
10 and
(3) a catalytic amount of an addition reaction catalyst.

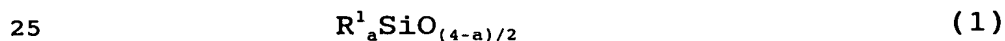
The organic peroxide curing type organopolysiloxane composition is preferably defined as comprising

- 15 (i) 100 parts by weight of an organopolysiloxane having on the average at least two alkenyl groups in a molecule, and
(ii) a catalytic amount of an organic peroxide.

[0010]

In the addition reaction curing type organopolysiloxane composition, the organopolysiloxane
20 having on the average at least two alkenyl groups in a molecule as component (1) is typically represented by the following average compositional formula (1):

[0011]



wherein R^1 , which may be the same or different, is a substituted or unsubstituted monovalent hydrocarbon group of 1 to 12 carbon atoms, preferably 1 to 8 carbon atoms, and
30 "a" is a positive number of 1.5 to 2.8, preferably 1.8 to 2.5, and more preferably 1.95 to 2.05.

[0012]

Examples of the substituted or unsubstituted monovalent hydrocarbon groups attached to silicon atoms,
35 represented by R^1 , include alkyl groups such as methyl, ethyl, propyl, isopropyl, butyl, isobutyl, tert-butyl, pentyl, neopentyl, hexyl, cyclohexyl, octyl, nonyl and decyl; aryl groups such as phenyl, tolyl, xylyl and

naphthyl; aralkyl groups such as benzyl, phenylethyl and phenylpropyl; alkenyl groups such as vinyl, allyl, propenyl, isopropenyl, butenyl, hexenyl, cyclohexenyl and octenyl; and substituted ones of the foregoing groups in which some or all of the hydrogen atoms are replaced by halogen atoms (e.g., fluoro, bromo and chloro), cyano groups or the like, such as chloromethyl, chloropropyl, bromoethyl, trifluoropropyl and cyanoethyl.

[0013]

At least two of the R^1 groups must be alkenyl groups, preferably of 2 to 8 carbon atoms, more preferably 2 to 6 carbon atoms. It is noted that the content of alkenyl groups is preferably 0.001 to 20 mol%, especially 0.01 to 10 mol% based on the entire organic groups attached to silicon atoms (that is, the substituted or unsubstituted monovalent hydrocarbon groups included as R^1 in formula (1)). The alkenyl groups may be attached to silicon atoms at ends of the molecular chain and/or silicon atoms midway the molecular chain. When the cure rate of the composition and the physical properties of cured parts are taken into account, the organopolysiloxane should preferably have at least alkenyl groups attached to the silicon atoms at molecular chain ends. The preferred groups represented by R^1 other than the alkenyl groups are alkyl groups such as methyl, aryl groups such as phenyl, and fluoroalkyl groups such as trifluoropropyl. Especially alkyl groups are preferred.

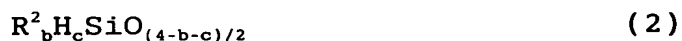
[0014]

With respect to the structure, the organopolysiloxane is generally a diorganopolysiloxane of a basically straight chain structure whose backbone is comprised of recurring diorganosiloxane units $R^1_2SiO_{2/2}$ and which is blocked with a triorganosiloxy group $R^1_3SiO_{1/2}$ at either end of the molecular chain. However, it may have a partially branched or cyclic structure containing $R^1SiO_{3/2}$ and/or $SiO_{4/2}$ units. The alkenyl group-containing organopolysiloxane may have any desired degree of polymerization or viscosity, and encompasses from

ones having a low degree of polymerization which are liquid at room temperature (25°C) to gum-like ones having a high degree of polymerization. Often, the organopolysiloxane used herein has an average degree of polymerization (weight average degree of polymerization) of 10 to 10,000, preferably 50 to 10,000, more preferably 100 to 10,000, and most preferably 100 to 2,000. With an average degree of polymerization of less than 10, the cured rubber sometimes has unsatisfactory physical properties. It is noted that the organopolysiloxane preferably has a viscosity of 100 to 1,000,000 cps (centipoise) at 25°C, and more preferably 500 to 500,000 cps at 25°C.

[0015]

As component (2), the organohydrogenpolysiloxane is typically represented by the following average compositional formula (2):



and should have at least two (generally 2 to 300), preferably at least three, and more preferably 3 to about 150, silicon atom-attached hydrogen atoms (Si-H groups).

[0016]

In formula (2), R^2 is a substituted or unsubstituted monovalent hydrocarbon group of 1 to 12 carbon atoms, examples of which are as described for R^1 in formula (1). Hydrocarbon groups free of aliphatic unsaturation are preferred, with alkyl groups such as methyl being especially preferred. The letter b is a positive number of 0.7 to 2.1, c is a positive number of 0.001 to 1, and b+c is 0.8 to 3. Preferably, b is 1 to 2, c is 0.01 to 1, and b+c is 1.5 to 2.8.

[0017]

At least two, preferably at least three Si-H groups in the molecule may be positioned at ends and/or midway of the molecular chain. The molecular structure of organohydrogenpolysiloxane may be straight, cyclic, branched or three-dimensional network. Desirable are those which are

liquid at room temperature (25°C) and have a number of silicon atoms per molecule or a degree of polymerization in the range of 2 to 300, more desirably 3 to 150. It is noted that the organohydrogenpolysiloxane preferably has a
5 viscosity of 0.1 to 10,000 cps at 25°C, and more preferably 0.5 to 5,000 cps at 25°C.

[0018]

Exemplary organohydrogenpolysiloxanes of formula (2) include 1,1,3,3-tetramethyldisiloxane, methylhydrogencyclo-
10 polysiloxane, methylhydrogensiloxane-dimethylsiloxane cyclic copolymers, both end trimethylsiloxy-blocked methylhydrogenpolysiloxane, both end trimethylsiloxy-blocked dimethylsiloxane-methylhydrogensiloxane copolymers, both end dimethylhydrogensiloxy-blocked dimethylpolysiloxane, both
15 end dimethylhydrogensiloxy-blocked dimethylsiloxane-methylhydrogensiloxane copolymers, both end trimethylsiloxy-blocked methylhydrogensiloxane-diphenylsiloxane copolymers, both end trimethylsiloxy-blocked methylhydrogensiloxane-diphenylsiloxane-dimethylsiloxane copolymers, both end
20 dimethylhydrogensiloxy-blocked methylhydrogensiloxane-dimethylsiloxane-diphenylsiloxane copolymers, copolymers of $(\text{CH}_3)_2\text{HSiO}_{1/2}$ units, $(\text{CH}_3)_3\text{SiO}_{1/2}$ units, and $\text{SiO}_{4/2}$ units, copolymers of $(\text{CH}_3)_2\text{HSiO}_{1/2}$ units and $\text{SiO}_{4/2}$ units, and copolymers of $(\text{CH}_3)_2\text{HSiO}_{1/2}$ units, $\text{SiO}_{4/2}$ units, and $(\text{C}_6\text{H}_5)_3\text{SiO}_{1/2}$
25 units.

[0019]

The organohydrogenpolysiloxane (2) is blended in an amount of 0.1 to 300 parts, preferably 0.3 to 200 parts and more preferably 0.5 to 100 parts by weight per 100 parts by
30 weight of the organopolysiloxane (1). Differently stated, the organohydrogenpolysiloxane (2) is blended in such an amount that 0.3 to 20 mol, preferably 0.5 to 5 mol, more preferably 0.8 to 3 mol, and most preferably 0.8 to 2.5 mol of silicon atom-attached hydrogen atoms (Si-H groups) in
35 component (2) are available per mol of silicon atom-attached alkenyl groups in component (1).

[0020]

Component (3) is an addition reaction catalyst. Platinum group metal catalysts are typical, including platinum catalysts, for example, platinum black, platinum chloride, chloroplatinic acid, reaction products of chloroplatinic acid with monohydric alcohols, complexes of chloroplatinic acid with olefins, and platinum bisacetoacetate, palladium catalysts, and rhodium catalysts. The addition reaction catalyst (3) is used in a catalytic amount, typically 0.1 to 1,000 ppm, preferably 0.5 to 1,000 ppm, and more preferably 1 to 500 ppm.

[0021]

The addition reaction curing type organopolysiloxane composition is comprised of components (1), (2) and (3) as mentioned above.

[0022]

The organopolysiloxane having on the average at least two alkenyl groups in a molecule used herein as component (i) may be the same as component (1) defined above.

[0023]

The organic peroxide (ii) used herein may be selected from conventional well-known ones, for example, benzoyl peroxide, 2,4-dichlorobenzoyl peroxide, p-methylbenzoyl peroxide, o-methylbenzoyl peroxide, 2,4-dicumyl peroxide, 2,5-dimethyl-bis(2,5-t-butylperoxy)hexane, di-t-butyl peroxide, t-butyl perbenzoate, 1,1-bis(t-butylperoxy)-3,3,5-trimethylcyclohexane, and 1,6-bis(t-butylperoxycarboxy)-hexane.

[0024]

The organic peroxide is used in a catalytic amount, usually 0.01 to 10 parts and preferably 0.1 to 10 parts by weight per 100 parts by weight of the organopolysiloxane (i).

[0025]

The organic peroxide curing type organopolysiloxane composition is comprised of components (i) and (ii) as mentioned above.

[0026]

The reinforcing silica fines (B) may be any of silica species commonly used in the art as a reinforcement for rubber.

5 [0027]

Although any of silica species used in conventional silicone rubber compositions is useful, reinforcing silica fines having a specific surface area of at least 50 m²/g are preferred. It is advantageous to use precipitated silica, fumed silica and fired silica having a specific surface area of 50 to 400 m²/g, with the fumed silica being especially advantageous for improved rubber strength. The reinforcing silica fines may be hydrophibized on surfaces with organosilicon compounds and other surface treating agents.

15 [0028]

It is recommended that silica fines be previously treated directly in powder form. Any of well-known techniques may be used for the surface treatment. For example, untreated silica fine powder and a surface treating agent are admitted into a mechanical milling device closed under atmospheric pressure or a fluidized bed where they are mixed to effect surface treatment at room temperature or elevated temperature, optionally in the presence of an inert gas. If desired, a catalyst is used to promote the surface treatment. After thorough admixing, the powder is dried. The amount of the surface treating agent used is at least the theoretical amount calculated from the surface area to be covered with the agent.

[0029]

30 Suitable treating agents include organosilazanes such as hexamethyldisilazane and divinyltetramethyldisilazane; silane coupling agents, for example, organoalkoxysilanes such as methyltrimethoxysilane, ethyltrimethoxysilane, propyltrimethoxysilane, butyltrimethoxysilane, 35 dimethyldimethoxysilane, diethyldimethoxysilane, vinyltriethoxysilane, vinyltrimethoxysilane, trimethylmethoxysilane, triethylmethoxysilane,

vinyltris(methoxyethoxy)silane, trimethylchlorosilane, dimethyldichlorosilane, divinyltrimethoxysilane and chloropropyltrimethoxysilane; and organosilicon compounds, for example, organopolysiloxanes such as

5 polydimethylsiloxane and organohydrogenpolysiloxane. After surface treatment with such treating agents, the resulting hydrophobic silica fines are ready for use. Of these treating agents, the silane coupling agents and silazanes are preferred.

10 [0030]

An appropriate amount of component (B) blended is 1 to 100 parts, more preferably 2 to 80 parts by weight per 100 parts by weight of component (A). Less amounts of component (B) fail to provide mechanical strength whereas excessive
15 amounts of component (B) are difficult to fill, aggravating working and processing efficiency.

[0031]

It is recommended that when the surface treated (hydrophobic) silica fines are admixed with component (A),
20 the silica fines be further subjected to a hydrophobic treatment. That is, a surface treating agent is used in the admixing step for imparting additional hydrophobic property to the once hydrophobized silica fines. Examples of the surface treating agent include silane coupling agents and
25 partial hydrolyzates thereof, organosilazanes, titanate coupling agents, organopolysiloxane fluids, and organohydrogenpolysiloxane fluids.

[0032]

The surface treating agents used herein are described
30 in further detail. Exemplary silane coupling agents used herein are organoalkoxysilanes including methyltrimethoxysilane, ethyltrimethoxysilane, propyltrimethoxysilane, butyltrimethoxysilane, dimethyldimethoxysilane, diethyldimethoxysilane,
35 vinyltriethoxysilane, vinyltrimethoxysilane, trimethylmethoxysilane, triethylmethoxysilane, vinyltris(methoxyethoxy)silane, trimethylchlorosilane,

trimethylaminosilane, glycidoxypropyltrimethoxysilane, glycidoxypropylmethyldiethoxysilane, (epoxycyclohexyl)ethyltrimethoxysilane, methacryloxypropyltrimethoxysilane, 5 methacryloxypropyltriethoxysilane, dimethyldimethoxysilane, divinylldimethoxysilane, and chloropropyltrimethoxysilane. Other silanes are also useful. Partial hydrolyzates of these silanes are also useful.

[0033]

10 Exemplary organosilazanes used herein are hexamethyldisilazane, divinyltetramethyldisilazane and diphenyltetramethyldisilazane.

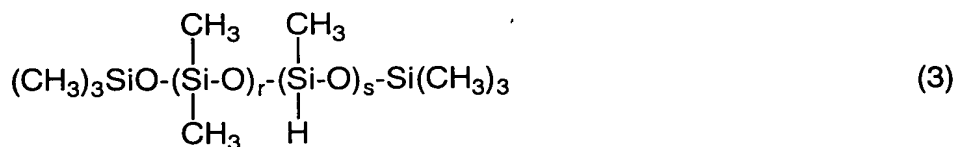
[0034]

Exemplary titanate coupling agents used herein include 15 tetraisopropyl titanate, tetra-n-butyl titanate, butyl titanate dimer, tetrastearyl titanate, triethanolamine titanate, titanium acetylacetonate, titanium ethylacetoacetate, titanium lactate, octylene glycol titanate, isopropyl tristearoyl titanate, isopropyl 20 tridodecylbenzenesulfonyl titanate, isopropyl tris(dioctylpyrophosphate) titanate, bis(dioctylpyrophosphate) oxyacetate titanate, and bis(dioctylpyrophosphate) ethylene titanate.

[0035]

25 The organopolysiloxane fluids used herein may be cyclic, linear, branched or network although linear or cyclic dimethylpolysiloxanes having a viscosity of 0.65 to 100,000 centistokes at 25°C are preferred. The organohydrogenpolysiloxane fluids used herein may have any 30 of cyclic, linear, branched and network molecular structures. Linear methylhydrogenpolysiloxanes of the following average formula (3) are desirably used.

[0036]



[0037]

In the formula, r is an integer of 0 to 50, and s is an integer of 1 to 50. Siloxanes with r in excess of 50 are too viscous and difficult to treat therewith. Similarly,
5 siloxanes with s in excess of 50 are too viscous and difficult to wet the surface.

[0038]

The amount of the surface treating agent used is at least the theoretical amount calculated from the surface
10 area to be covered with the agent although amounts less than the theory are acceptable if no problems arise. Specifically, less than 0.1 parts by weight of the second agent per 100 parts by weight of the silica fines is ineffective whereas more than 20 parts by weight of the
15 second agent is wasteful and adds to the cost. For this reason, 0.1 to 20 parts, especially 0.5 to 10 parts by weight of the second surface treating agent is preferably used per 100 parts by weight of the silica fines.

[0039]

20 The surface treatment may be effected on the silica fines as an independent step or together with the step of mixing with other components. Any of well-known techniques for surface treatment may be used. For example, the treated silica fine powder, other components and the second surface
25 treating agent are admitted into a mechanical milling device under atmospheric pressure where they are mixed to effect surface treatment at room temperature or elevated temperature, optionally in the presence of an inert gas. If desired, a catalyst is used to promote the surface
30 treatment.

[0040]

An organic compound or organosilicon compound having an epoxy equivalent of 100 to 5,000 g/mol and containing at least one aromatic ring in a molecule is component (C)
35 essential for imparting adhesion, also referred to as tackifier component.

[0041]

It should contain at least one, preferably 1 to 20, aromatic ring (e.g., benzene ring, naphthalene ring, anthracene ring) in a molecule. The inclusion of an aromatic ring in the molecule reduces compatibility with component (A) and improves adhesion at the rubber/resin interface. If component (C) is compatible with component (A), it is likely that more of this tackifier component is localized within the rubber component, adversely affecting adhesion to resins. It is thus recommended that component (C) be semi-compatible or incompatible with component (A).

[0042]

The organic or organosilicon compound should also have at least one epoxy group (e.g., glycidoxy) in the molecule. The epoxy group such as glycidoxy is necessary for the compound to develop adhesion. The epoxy equivalent is in the range of 100 to 5,000 g/mol, and preferably 120 to 4,000 g/mol. An epoxy equivalent below 100 g/mol leads to difficulty of synthesis, too poor compatibility with the silicone rubber composition, and detrimental effects on rubber physical properties. An epoxy equivalent in excess of 5,000 g/mol leads to insufficient adhesion.

[0043]

As component (C), compounds containing at least one hydrogen atom attached to a silicon atom (i.e., Si-H group) in the molecule are preferred. Often organosilicon compounds, such as organosilanes and organosiloxanes, having at least one Si-H group, usually 1 to 100, preferably 1 to 50, more preferably 2 to 20 Si-H groups in the molecule, and having 1 to 500, preferably 1 to 200 silicon atoms are used.

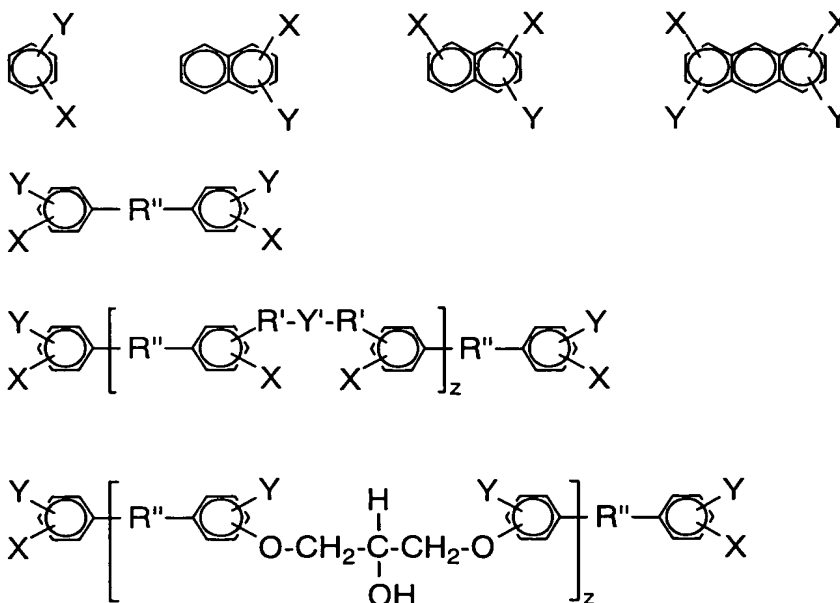
[0044]

As mentioned above, component (C) used herein is an organic compound or organosilicon compound (such as organosilane and organosiloxane) containing an amount of epoxy group to give an epoxy equivalent of 100 to 5,000 g/mol, and having a phenyl skeleton or at least one benzene ring (as typified by phenyl group), naphthalene ring or

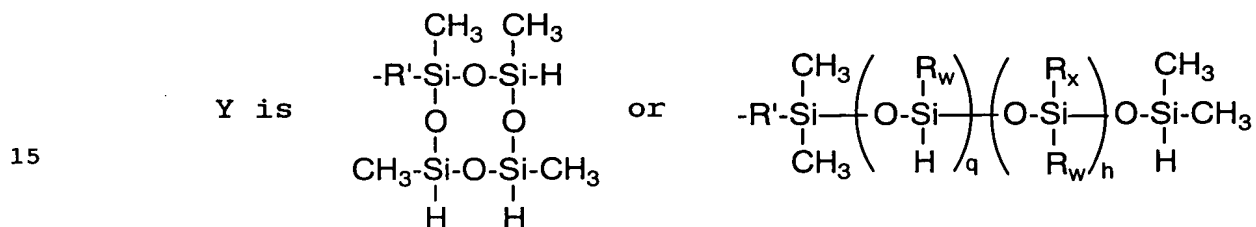
anthracene ring as the aromatic ring in the molecule; and preferably an organosilicon compound containing an amount of epoxy group to give an epoxy equivalent of 100 to 5,000 g/mol, having a phenyl skeleton or at least one benzene ring (as typified by phenyl group), and having at least one Si-H group in the molecule. It is noted that JP-A 6-172738 and JP-A 8-53661 disclose an adhesive silicone rubber composition and a silicone adhesive composition, respectively, which contains a tackifier component having a phenyl skeleton or benzene ring (as typified by phenyl group) in the molecule, but not an epoxy group. These adhesive silicone rubber compositions are insufficiently adherent to thermoplastic resins such as polymethyl methacrylate, polyphenylene sulfide, polyamide and polyphthalamide. Separately, compositions comprising a tackifier component having Si-H groups and epoxy groups are known in the art, but this tackifier component does not include a phenyl skeleton or a benzene ring structure (as typified by phenyl group) in the molecule. These compositions fail to achieve sufficient adhesion to the aforementioned thermoplastic resins as well.

[0045]

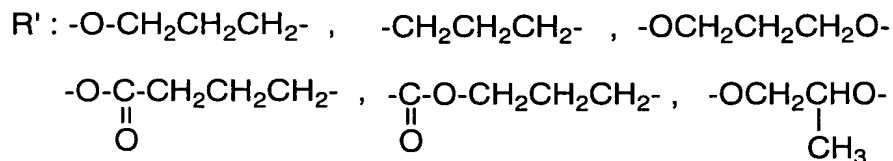
Illustrative examples of the compound (C) used herein are given below.



In the formulae, X is $\begin{array}{c} -O-CH_2-CH-CH_2 \\ \quad \quad \quad \diagup \quad \diagdown \\ \quad \quad \quad O \end{array}$.



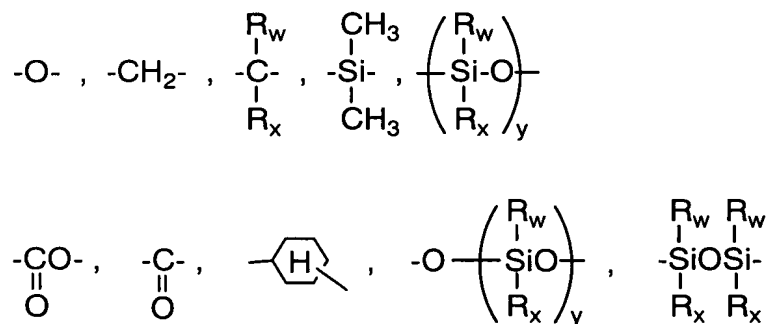
20 wherein R' is selected from the following groups:



Rw and Rx are substituted or unsubstituted monovalent hydrocarbon groups, q is a number of 1 to 50, preferably 1 to 20, and h is a number of 0 to 50, preferably 1 to 100.

25

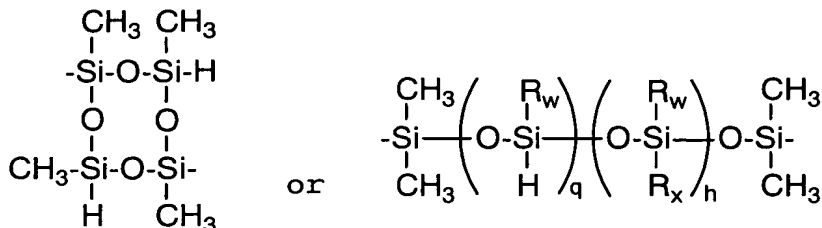
R" is selected from the following groups:



wherein Rw and Rx are as defined above, and y is a number of 0 to 100.

30

Y' is



wherein R_w , R_x , q and h are as defined above. Subscript z is a number of 1 to 10.

5 [0046]

Also useful are the organic compounds or organosilicon compounds obtained by incorporating alkoxysilyl groups (e.g., trimethoxysilyl, triethoxysilyl or methyldimethoxysilyl), acrylic, methacrylic, ester, carboxyl
10 anhydride, amino or amide groups in the above-enumerated compounds.

[0047]

More specifically, the substituted or unsubstituted monovalent hydrocarbon groups represented by R_w and R_x are
15 preferably those of 1 to 12 carbon atoms, especially 1 to 8 carbon atoms. Examples include alkyl, aryl, aralkyl, alkenyl and other groups as enumerated for R^1 . Exemplary substituted monovalent hydrocarbon groups are those substituted with alkoxy, acrylic, methacrylic, acryloyl,
20 methacryloyl, amino and alkylamino groups.

[0048]

As the tackifier component, the above-enumerated compounds may be used alone or in admixture. The reaction product of the above-enumerated compounds is also useful.

25 [0049]

An appropriate amount of the tackifier component (C) blended is 0.1 to 50 parts by weight per 100 parts by weight of the organopolysiloxane composition (A). With less than 0.1 part of component (C), sufficient adhesion is not
30 accomplished. More than 50 parts of component (C) can

degrade physical properties. The preferred amount is 0.2 to 30 parts by weight.

[0050]

For the purpose of effectively assisting the tackifier
5 component (C), an organic compound or organosilicon compound
having one alkenyl group and at least one ester group in a
molecule may be blended. Examples are organic compounds or
organosilicon compounds having incorporated therein
unsaturated carboxylic acids such as acrylic acid,
10 methacrylic acid and vinylacetic acid, and allyl esters such
as allyl benzoate, diallyl phthalate, tetraallyl
pyromellitate, and alkyl acid allyl esters. It is also
acceptable to add epoxy ring-opening catalysts such as
organometallic chelates, amines, amides, imidazoles and acid
15 anhydrides. These aids are added in such amounts that the
benefits of the invention are not impaired.

[0051]

Besides, a siloxane skeleton-bearing organosilicon
compound incompatible with component (A) may be added for
20 the purpose of improving adhesion. Examples include phenyl
group-bearing silicone oil, fluorosilicone oil,
silicone-modified perfluoroalkyl ether compounds,
polyether-modified silicone oil, amino-modified silicone
oil, amide-modified silicone oil, imide-modified silicone
25 oil, and urethane-modified silicone oil.

[0052]

To impart an oil bleeding ability to the inventive
composition, at least one non-functional organopolysiloxane
having phenyl groups may be added in an amount of about 1 to
30 20 parts per 100 parts by weight of component (A).

[0053]

In addition to the above-described components, various
additives may be added to the silicone rubber adhesive
composition of the invention. Such additives are selected
35 for a particular purpose. Typical additives are metal
oxides and compounds thereof such as titanium oxide, iron
oxide, cerium oxide, vanadium oxide, cobalt oxide, chromium

oxide, and manganese oxide, and inorganic fillers, for example, quartz powder, diatomaceous earth, calcium carbonate, magnesium carbonate, alumina, and carbon. Also, pigments, heat resistance modifiers, flame retardants, plasticizers, and reaction regulators may be added insofar as the desired properties are not impaired. These optional additives may be added in conventional amounts that would not hamper the benefits of the invention.

[0054]

In the embodiment wherein the addition curing type organopolysiloxane composition is used as (A), an addition reaction regulator may be added insofar as the objects of the invention are not impaired. Suitable regulators are vinyl group-bearing organopolysiloxanes such as vinylcyclotetrasiloxane, triallyl isocyanate, alkyl maleates, acetylene alcohols such as ethynyl cyclohexanol, silanes, modified siloxanes, hydroperoxides, tetramethylethylenediamine, benzotriazole and mixtures of any.

[0055]

The silicone rubber adhesive composition of the invention can be obtained simply by uniformly admixing the above-described components (A) to (C) and optional components at room temperature. Preferably, component (B) is mixed with component (A) excluding (2) and (3) or (ii) in a planetary mixer or kneader where they are heat treated at a temperature of 100 to 200°C for 1 to 4 hours, during which period the second surface treating agent may be added. Thereafter, at room temperature, the mixture is admixed with components (2) and (3) or (ii) of component (A) and component (C), thereby obtaining the composition. Any desired molding technique may be selected depending on the viscosity of the mixture or composition. Any of casting, compression molding, injection molding, extrusion molding, and transfer molding techniques is useful. The composition may be heat molded, typically at a temperature of about 60 to 200°C for about 10 seconds to about 24 hours.

[0056]

The silicone rubber adhesive composition of the invention is advantageously used in forming integrally molded articles with organic resins. The organic resins
5 used in combination with the silicone rubber adhesive composition are thermoplastic resins of polymerized or polycondensed olefin family, for example, acrylonitrilebutadiene-styrene (ABS) resins, styrene resins, polyethylene resins, polypropylene resins, polyacetal
10 resins, acrylic resins, polycarbonate (PC) resins, polyethylene terephthalate resins, polybutylene terephthalate (PBT) resins, polyphenylene oxide resins, phenolic resins, epoxy resins, polyphenylene sulfide resins, polysulfone resins, nylon resins, aromatic polyamide resins,
15 polyimide resins, liquid crystal resins, and mixtures thereof. They may also take the form of fibers or film.

[0057]

Several methods may be employed in integrally molding the uncured silicone rubber adhesive composition on the
20 thermoplastic resin. In one method, a desired shape of the uncured silicone rubber adhesive composition is placed on a preform of the thermoplastic resin, followed by heating at a temperature below the melting temperature of the thermoplastic resin. Another method is by placing the
25 uncured silicone rubber adhesive composition on a preform of the thermoplastic resin, followed by compression at a temperature below the melting temperature of the thermoplastic resin. A further method is by previously injection molding the thermoplastic resin in a mold by means
30 of an injection molding machine, and heat injecting the silicone rubber adhesive composition into the mold. It is noted that the preform of the thermoplastic resin is conveniently prepared by a well-known method involving pelletizing the thermoplastic resin, injecting the pellets
35 into a mold heated above the softening point of the thermoplastic resin, then cooling the mold below the softening point of the thermoplastic resin. In these

molding steps, customary molding machines such as injection molding machines and transfer molding machines can be employed. The mold used herein may be a mold of a metal such as steel or stainless steel which has not been surface
5 treated, or a metal mold which has been surface treated such as by chromium plating or nickel plating.

[0058]

The silicone rubber adhesive composition in the uncured state may take the form of liquid, putty or paste,
10 with the liquid or paste form being preferred for ease of molding. In order that the silicone rubber adhesive composition firmly bond with the thermoplastic resin, the conditions for curing the silicone rubber adhesive composition should include a temperature and time which do
15 not cause deformation, melting and degradation of the thermoplastic resin. Usually integrally molded articles can be formed under curing conditions of 50 to 150°C and about 0.2 to 5 hours, especially about 0.4 to 30 minutes, although suitable curing conditions vary with the type of resin.

20 [0059]

[Effect of the Invention]

In connection with an integrally molded article of a silicone rubber adhesive composition and a thermoplastic resin, the silicone rubber adhesive composition of the
25 invention is easily moldable within a short time by injection molding, suitable in primerless molding, and bondable with the thermoplastic resin. A bond between the silicone rubber and the thermoplastic resin is achieved without a need for modification of the resin. The
30 integrally molded articles are useful in the electric, electronic, automotive, precision machinery and building fields.

[0060]

[EXAMPLE]

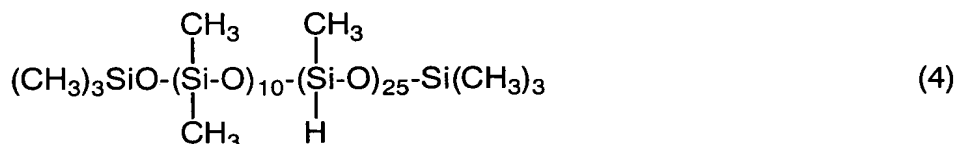
35 Examples and Comparative Examples of the invention are given below by way of illustration and not by way of limitation. All parts are by weight.

Preparation examples of silicone rubber adhesive compositions

As shown in Table 1, a dimethylpolysiloxane blocked with a dimethylvinylsiloxyl group at each end and having a viscosity of 10,000 cps at 25°C as component (1) of component (A) and fumed silica having a specific surface area of 200 m²/g as component (B) were blended together with 8 parts per 100 parts of component (1) of hexamethyldisilazane and uniformly admixed in a kneader/mixer, and further heat admixed at 150°C for one hour, obtaining a silicone rubber base. To the silicone rubber base were added methylhydrogenpolysiloxane of the average compositional formula (4) as component (2) in component (A), a 1% 2-ethylhexanol solution of chloroplatinic acid as component (3), and a 50% ethanol solution of ethynyl cyclohexanol as a reaction regulator. Further, tackifier I to IV as component (C) was added. They were uniformly mixed, yielding a silicone rubber adhesive composition.

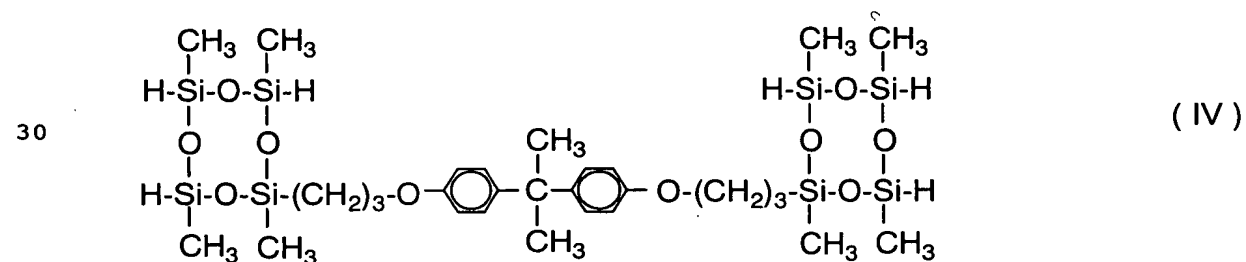
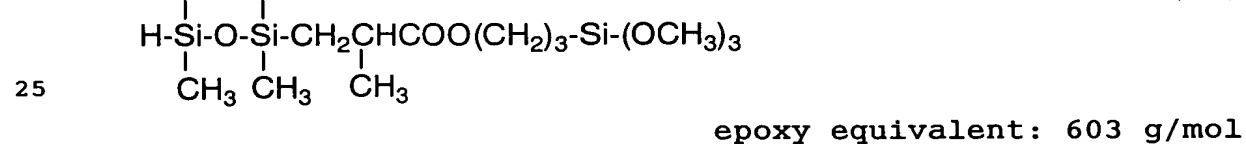
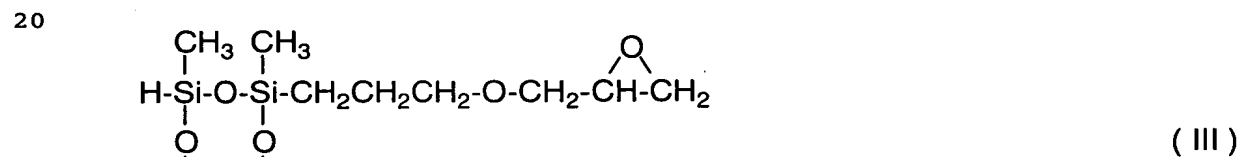
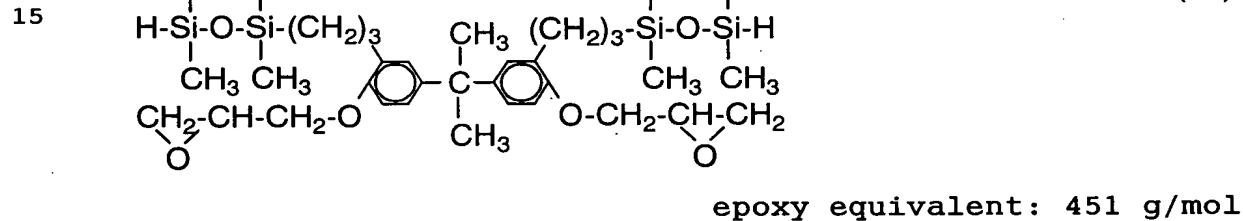
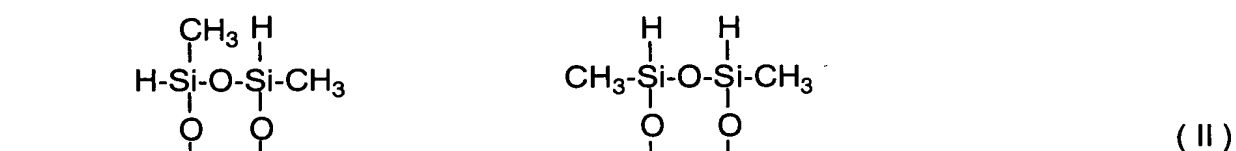
[0061]

Component (2)



[0062]

Component (C)



epoxy equivalent: 0 g/mol

[0063]

[Example 1]

A thermoplastic resin injection molding machine was used. A polycarbonate (PC) resin having a number average molecular weight of about 10,000, polybutylene terephthalate (PBT) resin having a number average molecular weight of about 20,000, polymethyl methacrylate (PMMA) resin, polyphenylene sulfide (PPS) resin, nylon 6-6 (PA66), and polyphthalamide (PPA) each were admitted into the machine, plasticized at 290°C and injected into a plurality of cavities in a multiple sheet mold, forming resin sheets of 25 mm × 100 mm × 2 mm (thick). The injection conditions included an injection time of 6 seconds, a cooling time of 30 seconds, an injection pressure of 1,000 kg/cm², a clamping pressure of 35 ton, and a cavity temperature of 100°C. Each resin sheet and a chromium-plated metal plate of the same size were secured to a tensile shear bond strength test specimen-forming jig. An appropriate amount of the silicone rubber adhesive composition obtained in the above Preparation Example using tackifier I as component (C) was flowed into the jig, which was heated for 10 minutes in a thermostat tank at 120°C for curing. A test specimen as shown in FIG. 1 was obtained in this way and examined for bond strength according to JIS K-6850.

[0064]

In FIG. 1, "a" and "b" designate resin sheets, "c" and "d" are holder clamps, and "e" is the cured silicone rubber adhesive composition.

[0065]

[Example 2]

The resins used in Example 1 each were admitted into the injection molding machine, plasticized at 290°C and injected into a plurality of cavities in the multiple sheet mold, forming resin sheets of 25 mm × 100 mm × 2 mm (thick). The injection conditions included an injection time of 6 seconds, a cooling time of 30 seconds, an injection pressure of 1,000 kg/cm², a clamping pressure of 35 ton, and a cavity

temperature of 100°C. Each resin sheet and a chromium-plated metal plate of the same size were secured to the tensile shear bond strength test specimen-forming jig. An appropriate amount of the silicone rubber adhesive composition obtained in the above Preparation Example using tackifier II as component (C) was flowed into the jig, which was heated for 10 minutes in a thermostat tank at 120°C for curing. A test specimen as shown in FIG. 1 was obtained in this way and examined for bond strength according to JIS K-6850.

[0066]

[Comparative Example 1]

The resins used in Example 1 each were admitted into the injection molding machine, plasticized at 290°C and injected into a plurality of cavities in the multiple sheet mold, forming resin sheets of 25 mm × 100 mm × 2 mm (thick). The injection conditions included an injection time of 6 seconds, a cooling time of 30 seconds, an injection pressure of 1,000 kg/cm², a clamping pressure of 35 ton, and a cavity temperature of 100°C. Each resin sheet and a chromium-plated metal plate of the same size were secured to the tensile shear bond strength test specimen-forming jig. An appropriate amount of the silicone rubber adhesive composition obtained in the above Preparation Example using tackifier III as component (C) was flowed into the jig, which was heated for 10 minutes in a thermostat tank at 120°C for curing. A test specimen as shown in FIG. 1 was obtained in this way and examined for bond strength according to JIS K-6850.

[0067]

[Comparative Example 2]

The resins used in Example 1 each were admitted into the injection molding machine, plasticized at 290°C and injected into a plurality of cavities in the multiple sheet mold, forming resin sheets of 25 mm × 100 mm × 2 mm (thick). The injection conditions included an injection time of 6

seconds, a cooling time of 30 seconds, an injection pressure of 1,000 kg/cm², a clamping pressure of 35 ton, and a cavity temperature of 100°C. Each resin sheet and a chromium-plated metal plate of the same size were secured to the tensile shear bond strength test specimen-forming jig. An appropriate amount of the silicone rubber adhesive composition obtained in the above Preparation Example using tackifier IV was flowed into the jig, which was heated for 10 minutes in a thermostat tank at 120°C for curing. A test specimen as shown in FIG. 1 was obtained in this way and examined for bond strength according to JIS K-6850.

[0068]

[Example 3]

Molding of the resin and the silicone rubber composition used in Example 2 was carried out using a two-color injection molding machine equipped with two injection cylinders. In this molding machine, nozzles 1 and 2 of the injection cylinders are connected to a mold as shown in FIG. 2. The nozzle 1 is aligned with a mold parting line whereas the nozzle 2 is designed to inject the material into a cavity from its center on the right side of the mold. The mold consists of a left mold half 3 and a right mold half 4. Two recesses are formed in opposed surfaces of the mold halves 3 and 4 so that the recesses may define two cavities 5 and 6 when the mold halves are mated together (See FIG. 2).

[0069]

Using this injection molding machine, each resin was melted at 290°C and injected into the first cavity 5 through the nozzle 1 to form a resin sheet 7 (see FIG. 3). The injection conditions included an injection time of 6 seconds, a cooling time of 35 seconds, and a temperature of 100°C for both the cavity 5 and the left mold half 3.

[0070]

Next, the right mold half 4 was moved apart to open the mold, the left mold half 3, with the molded resin sheet 7 held in its recess, was rotated 180°, and the right mold

half 4 was moved back. The mold halves were clamped again. At this point, a cavity for molding a silicone rubber sheet was defined between the (lower) recess in the right mold half 4 and the opposing surface of the molded resin sheet 7 as shown in FIG. 4.

[0071]

In this state, the silicone rubber adhesive composition used in Example 1 (whose recipe is shown in Table 1) was injected from the nozzle 2 against the surface of the molded resin sheet 7, forming a rubber sheet 8 (see FIG. 5). The injection conditions included an injection time of 6 seconds, a cooling time of 90 seconds, a temperature of 100°C for the left mold half 3, and a temperature of 120°C for the right mold half 4.

[0072]

In this way, there was obtained a composite body of the resin sheet 7 bonded to the rubber sheet 8 as shown in FIG. 6. It is noted that both resin and rubber sheets were dimensioned 2.5 mm × 150 mm × 2 mm (thick). Mold release, dimensional precision and productivity were satisfactory. Adhesion was also examined.

[0073]

The test results are shown in Table 1.
Adhesive test condition:

It is noted that the bond strength was examined by a tensile shear bond strength test according to JIS K-6850. Specimens showing a bond strength of 25 kgf/cm² or higher were rated "Bonded."

[0074]

Table 1

	Example 1	Example 2	Example 3	Comparative Example 1	Comparative Example 2
(A) (1) Dimethylpolysiloxane	100	100	100	100	100
(2) Methylhydrogenpolysiloxane	5	5	5	5	5
(3) Platinum catalyst	0.1	0.1	0.1	0.1	0.1
Reaction regulator	0.05	0.05	0.05	0.05	0.05
(B) Silica fines	40	40	40	40	40
(C) Tackifier I	5				
II		5	5		
III				5	
IV					5
Adhesion Resin PC	Bonded	Bonded	Bonded	No	Bonded
PBT	Bonded	Bonded	Bonded	No	Bonded
PMMA	Bonded	Bonded	Bonded	No	No
PPS	Bonded	Bonded	Bonded	No	No
PA66	Bonded	Bonded	Bonded	No	No
PPA	Bonded	Bonded	Bonded	No	No
Metal (mold) Cr-plated	No	No	No	No	No

5 [0075]

It is evident from Table 1 that silicone rubber adhesive compositions within the scope of the invention can be briefly cured to various resins to form integrally molded articles in which the silicone rubber is firmly bonded to the resin. Additionally mold release, dimensional precision and productivity are satisfactory.

[BRIEF DESCRIPTION OF THE DRAWINGS]

[FIG. 1]

It is illustrates a bond strength test specimen, (A) being a cross-sectional view and (B) being a plan view.

[FIG. 2]

A schematic view of a two-color injection molding machine.

[FIG. 3]

5 A view similar to FIG. 2, showing the resin injected into a first cavity.

[FIG. 4]

A view similar to FIG. 2, showing the molded resin sheet set in a second cavity.

10 [FIG. 5]

A view similar to FIG. 2, showing the silicone rubber composition injected into the second cavity.

[FIG. 6]

15 A perspective view of the resulting composite body of resin and rubber sheets.

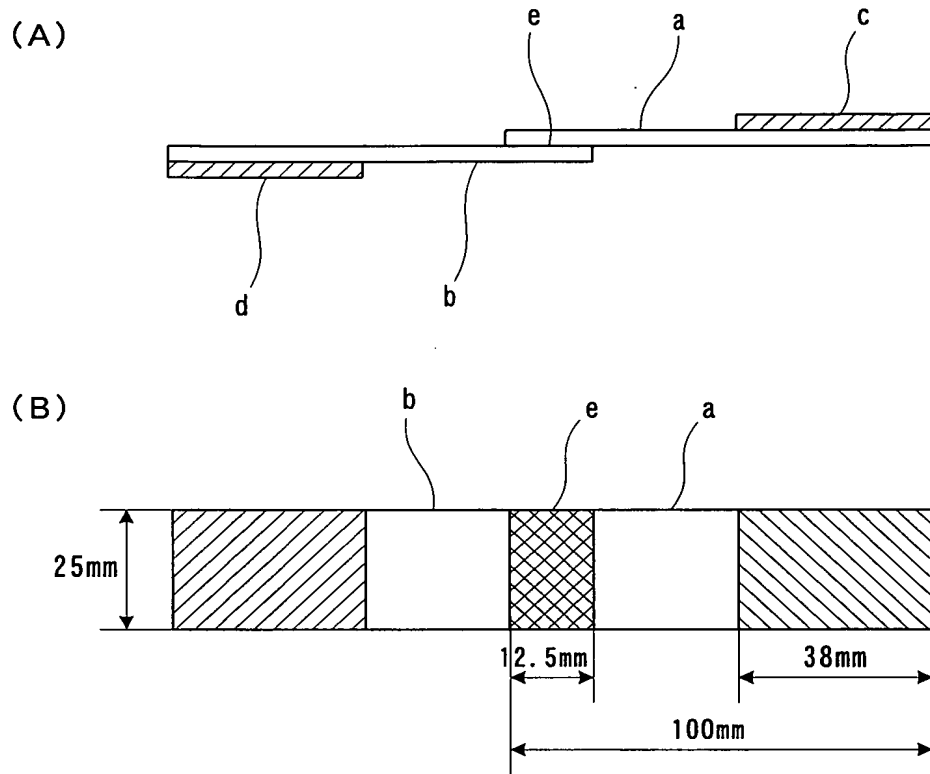
[Explanation of Symbols]

1	Nozzle
2	Nozzle
3	Left mold half
20 4	Right mold half
5	Cavity
6	Cavity
7	Resin sheet
8	Rubber sheet

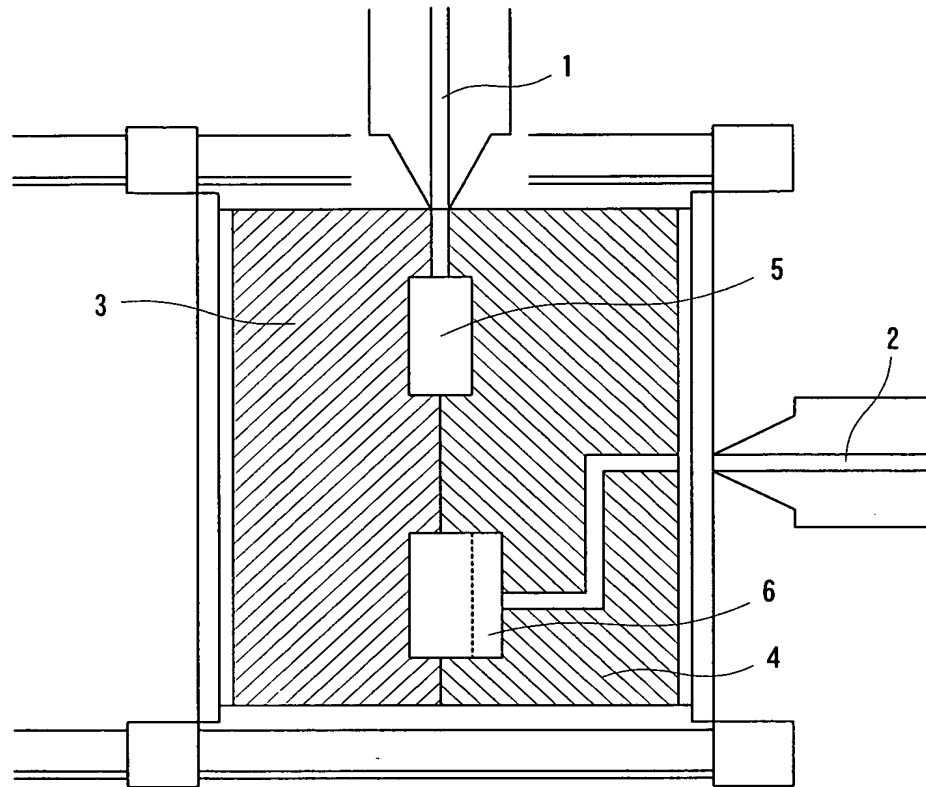
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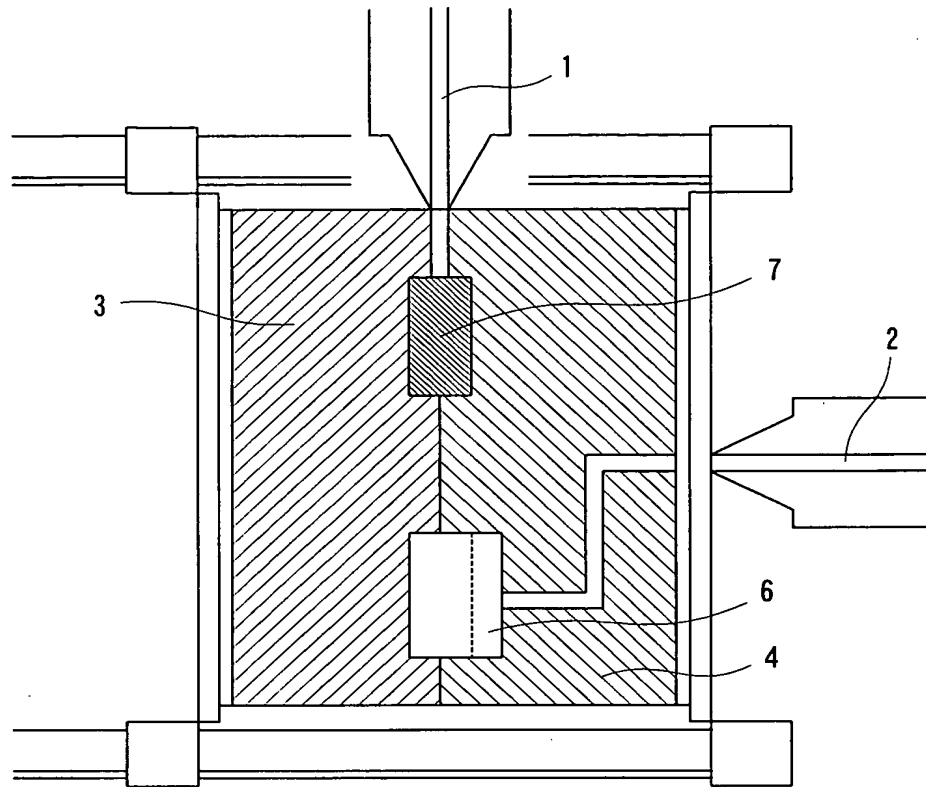
[FIG. 1]



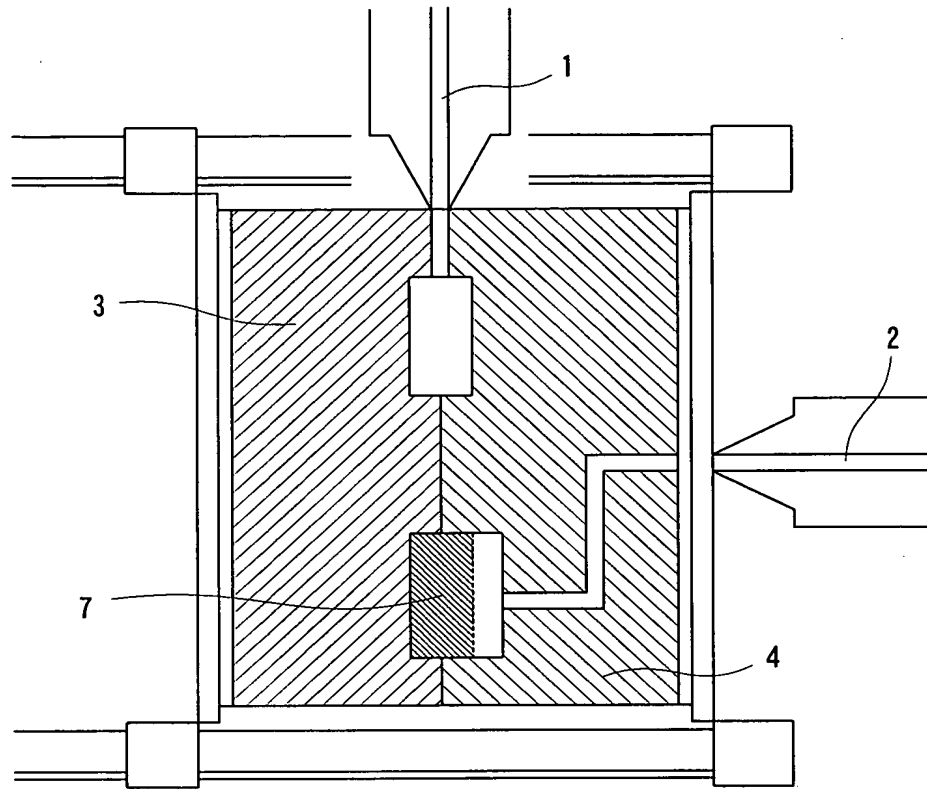
[FIG. 2]



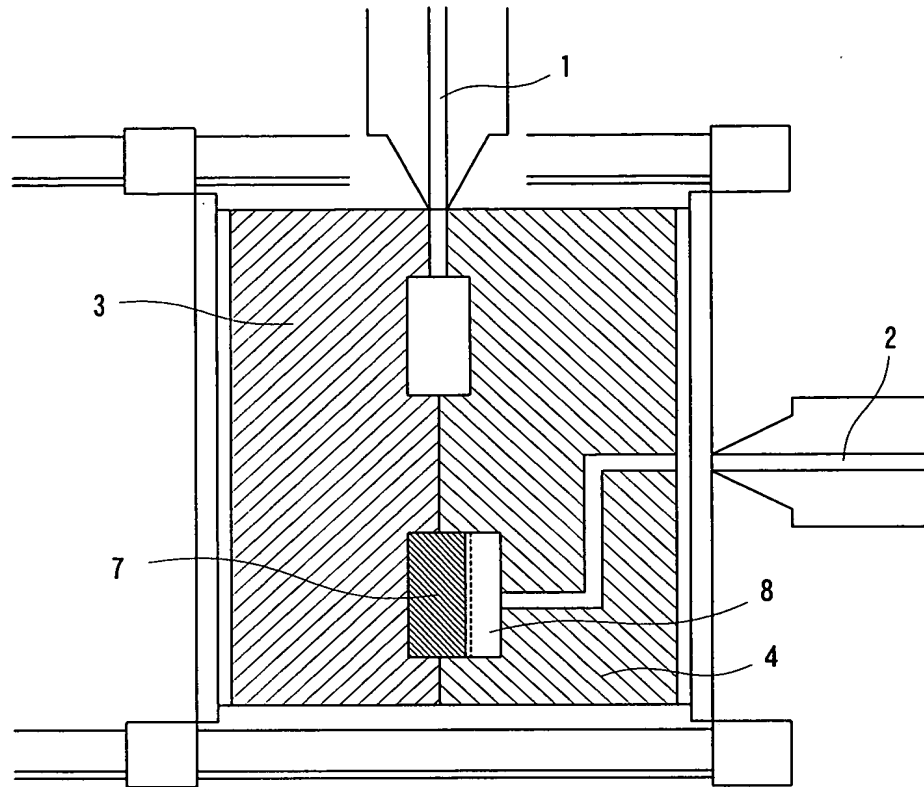
[FIG. 3]



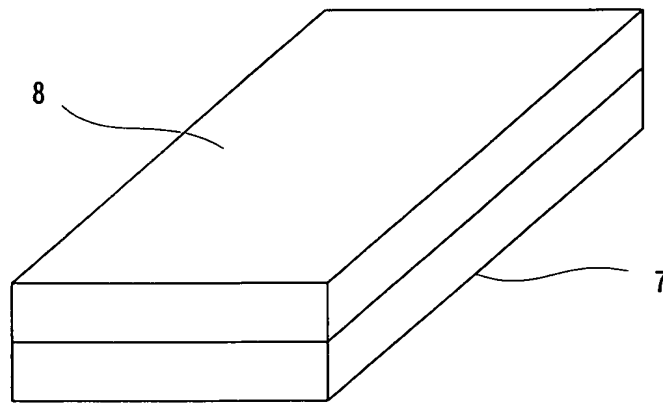
[FIG. 4]



[FIG. 5]



[FIG. 6]



2001-000536

[ABSTRACT]

[Means for Solution]

A silicone rubber adhesive composition comprising

5 (A) 100 parts by weight of a heat curable organopolysiloxane composition,

(B) 1 to 100 parts by weight of reinforcing silica fines, and

10 (C) 0.1 to 50 parts by weight of an organic compound or organosilicon compound having an epoxy equivalent of 100 to 5,000 g/mol and containing at least one aromatic ring in a molecule.

[Effect]

In connection with an integrally molded article of a silicone rubber adhesive composition and a thermoplastic resin, the silicone
15 rubber adhesive composition of the invention is easily moldable within a short time by injection molding, suitable in primerless molding, and bondable with the thermoplastic resin. A bond between the silicone rubber and the thermoplastic resin is achieved without a need for modification of the resin. The integrally molded
20 articles are useful in the electric, electronic, automotive, precision machinery and building fields.

[Selected Drawing] none

IN THE U.S. PATENT AND TRADEMARK OFFICE

APPLICANT: Syuuichi AZECHI et al.
SERIAL NO.: 10/035,213
FILED: January 4, 2002
FOR: Silicone Rubber Adhesive Composition
and Integrally Molded Article of
Silicone Rubber and Thermoplastic Resin
GROUP: 1712
EXAMINER: FEELY, MICHAEL J

D E C L A R A T I O N

Honorable Commissioner of Patents and Trademarks
Washington, D.C. 20231

Sir,

I, Syuuichi AZECHI, resident of c/o
Silicone-Electronics Materials Research Center,
Shin-Etsu Chemical Co., Ltd., 1-10, Oaza Hitomi,
Matsuida-machi, Usui-gun, Gunma-ken, Japan do hereby
declare that:



1. I was graduated from Master Course of Department of Engineering, Gunma University, Japan in March 1988. Since April 1988, I have been employed by Shin-Etsu Chemical Co., Ltd., the assignee of the above-identified application. I have been engaged in research and development relating to silicones in the laboratory of the Company.

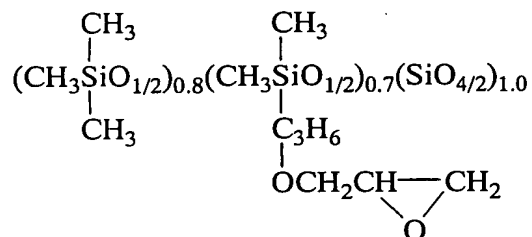
2. I am one of the named inventors of the above-identified application and hence, am familiar with the subject matter disclosed in said application.

3. In order to show the feature of the present invention, I conducted the following experiments.

[Experiment]

Comparison

Example 1 described in the specification of the above-identified application was repeated except that 5 parts by weight of an epoxy group-containing MQ resin:



disclosed in Preparation Example 1 of Morita '075 was used instead of tackifier I. The adhesion examined in the same manner as in the specification of the above-identified application. The results are shown in the following table. As is evident from the results, the effect of the present invention is not attained even if the above MQ resin is used.

Adhesion	Resin	PC	No
		PBT	No
		PMMA	No
		PPS	No
		PA66	No
		PPA	No
	Metal (mold)	Cr-plated	Bonded

I hereby declare that all statements made herein of my own knowledge are true and that all statements made on information and belief are believed to be true; and further that these statements were made with the knowledge that willful false statements and the like so made are punishable by fine or imprisonment, or both, under Section 1001 of Title 18 of the United States Code and that such willful false statements may jeopardize the validity of the application or any patent issued thereon.

Dated this 30th day of May, 2003

Symichi Azechi